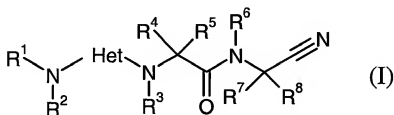


Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1. (Previously Presented) A compound of formula (I):



R<sup>1</sup> is independently hydrogen, C<sub>1-6</sub> alkyl or C<sub>3-6</sub> cycloalkyl;

R<sup>2</sup> is independently aryl, heteroaryl or a group C<sub>1-6</sub> alkylR<sup>9</sup>, CO(C<sub>1-6</sub> alkyl)R<sup>9</sup> or SO<sub>2</sub>(C<sub>1-6</sub>alkyl)R<sup>9</sup>; where R<sup>9</sup> is aryl or heteroaryl;

or R<sup>1</sup> and R<sup>2</sup> together with the nitrogen atom to which they are attached form a 4 to 7-membered saturated ring optionally containing a carbonyl group, O, S or N atom and optionally substituted by one or more C<sub>1-6</sub> alkyl, amino, hydroxy, CO<sub>2</sub>C<sub>1-6</sub> alkyl, COC<sub>1-6</sub> alkyl, halogen, C<sub>1-6</sub> alkylhydroxy, NR<sup>10</sup>R<sup>11</sup> where R<sup>10</sup> and R<sup>11</sup> are independently hydrogen, C<sub>1-6</sub> alkyl or together with the nitrogen atom to which they are attached form a 5- or 6-membered saturated ring optionally containing a further O, S or NR<sup>1</sup> group, C<sub>1-6</sub> alkylNR<sup>12</sup>R<sup>13</sup> where R<sup>12</sup> and R<sup>13</sup> are independently hydrogen or C<sub>1-6</sub> alkyl, CONR<sup>12</sup>R<sup>13</sup>, or optionally substituted by C<sub>1-6</sub> alkylR<sup>9</sup>, aryl, phenoxy, COaryl, COheteroaryl or a heteroaryl group, the latter six groups being optionally substituted by halogen, amino, hydroxy, cyano, nitro, carboxy, CONR<sup>12</sup>R<sup>13</sup>, SO<sub>2</sub>NR<sup>12</sup>R<sup>13</sup>, SO<sub>2</sub>R<sup>12</sup>, trifluoromethyl, NHSO<sub>2</sub>R<sup>12</sup>, NHCOR<sup>12</sup>, ethylenedioxy, methylenedioxy, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> alkyl NR<sup>10</sup>R<sup>11</sup>, SR<sup>12</sup> or NR<sup>10</sup>R<sup>11</sup>;

Het is a heteroaryl ring chosen from pyridine, pyrimidine, pyrazine, pyridazine or triazine and optionally substituted by halogen, amino, hydroxy, cyano, nitro, carboxy, CONR<sup>12</sup>R<sup>13</sup>, SO<sub>2</sub>NR<sup>12</sup>R<sup>13</sup>, SO<sub>2</sub>R<sup>12</sup>, trifluoromethyl, NHSO<sub>2</sub>R<sup>12</sup>, NHCOR<sup>12</sup>, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy, SR<sup>12</sup> or NR<sup>10</sup>R<sup>11</sup>;

R<sup>3</sup> is independently hydrogen, C<sub>1-6</sub> alkyl or C<sub>3-6</sub> cycloalkyl;

R<sup>4</sup> is independently hydrogen, C<sub>1-8</sub> alkyl, C<sub>3-8</sub> cycloalkyl, arylC<sub>1-5</sub>alkyl or heteroarylC<sub>1-5</sub>alkyl, the latter three groups being optionally substituted by one or more halogen, amino, hydroxy, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy, SR<sup>12</sup> or NR<sup>10</sup>R<sup>11</sup>;

R<sup>5</sup> is independently hydrogen, C<sub>1-6</sub> alkyl or C<sub>3-6</sub> cycloalkyl;

R<sup>6</sup> is independently hydrogen, C<sub>1-6</sub> alkyl or C<sub>3-6</sub> cycloalkyl;

R<sup>7</sup> is independently hydrogen, C<sub>1-6</sub> alkyl or C<sub>3-6</sub> cycloalkyl; and

R<sup>8</sup> is independently hydrogen, aryl, heteroaryl or C<sub>1-6</sub> alkyl optionally substituted with one or more aryl, heteroaryl, halogen, amino, hydroxy, carboxy, CONR<sup>12</sup>R<sup>13</sup>, SO<sub>2</sub>NR<sup>12</sup>R<sup>13</sup>, SO<sub>2</sub>R<sup>12</sup>, NHSO<sub>2</sub>R<sup>12</sup>, NHCOR<sup>12</sup>, C<sub>1-6</sub> alkyl, C<sub>3-6</sub> cycloalkyl, C<sub>1-6</sub> alkoxy, SR<sup>12</sup> or NR<sup>10</sup>R<sup>11</sup>;  
or a pharmaceutically acceptable salt thereof.

2. (Withdrawn) A compound according to claim 1 in which R<sup>1</sup> is hydrogen or C<sub>1-6</sub> alkyl and R<sup>2</sup> is CH<sub>2</sub>R<sup>9</sup> or CH<sub>2</sub>CH<sub>2</sub>R<sup>9</sup> where R<sup>9</sup> is phenyl or a 5- or 6-membered aromatic ring containing one or two heteroatoms and optionally substituted by C<sub>1-6</sub> alkyl.

3. (Previously Presented) A compound according to claim 1 in which R<sup>1</sup> and R<sup>2</sup> together with the nitrogen atom to which they are attached form a piperidine, piperazine, pyrrolidine, morpholine, or thiomorpholine ring optionally substituted by CH<sub>2</sub>OH, CH<sub>2</sub>CH<sub>2</sub>OH, hydroxy, CONH<sub>2</sub>, phenyl, phenoxy, or C(O)-furyl, the latter three groups being optionally substituted by halogen, in particular chloro.

4. (Previously Presented) A compound according to claim 1 in which R<sup>3</sup> is hydrogen.

5. (Previously Presented) A compound according to claim 1 in which R<sup>4</sup> is hydrogen.

6. (Previously Presented) A compound according to claim 1 in which R<sup>5</sup> is hydrogen or phenyl optionally substituted by C<sub>1-6</sub> alkyl or C<sub>1-6</sub> alkoxy.

7. (Currently Amended) A compound of formula (I) selected from:
- N~1~[Cyano(2-methoxyphenyl)methyl]-N~2~-(2-morpholin-4-ylpyrimidin-4-yl)-L-leucinamide,
- N~1~[Cyano(2-methoxyphenyl)methyl]-N~2~-(2-piperazin-1-ylpyrimidin-4-yl)-L-leucinamide,
- N-[Cyano(2-methoxyphenyl)methyl]-N-(2-morpholin-4-ylpyrimidin-4-yl)-L-phenylalaninamide,
- N~1~[Cyano(2-methoxyphenyl)methyl]-3-cyclohexyl-N~2~-(2-morpholin-4-ylpyrimidin-4-yl)-L-alaninamide,
- N-[2-(Benzylamino)pyrimidin-4-yl]-N-(cyanomethyl)-L-phenylalaninamide,
- N-{2-[Benzyl(methyl)amino]pyrimidin-4-yl}-N-(cyanomethyl)-L-phenylalaninamide,
- N-{2-[4-(4-Chlorophenyl)piperazin-1-yl]pyrimidin-4-yl}-N-(cyanomethyl)-L-phenylalaninamide,
- N~2~-[2-(Benzylamino)pyrimidin-4-yl]-N~1~-(cyanomethyl)-3-cyclohexyl-L-alaninamide,
- N~2~{2-[Benzyl(methyl)amino]pyrimidin-4-yl}-N~1~-(cyanomethyl)-3-cyclohexyl-L-alaninamide,
- N~2~{2-[4-(4-Chlorophenyl)piperazin-1-yl]pyrimidin-4-yl}-N~1~-(cyanomethyl)-3-cyclohexyl-L-alaninamide,
- N~1~-(Cyanomethyl)-N~2~-(4-morpholin-4-ylpyrimidin-2-yl)-L-leucinamide,
- N~1~-(Cyanomethyl)-N~2~-(2-morpholin-4-ylpyrimidin-4-yl)-L-leucinamide,
- N~1~-(Cyanomethyl)-N~2~-[2-(4-hydroxy-4-phenylpiperidin-1-yl)pyrimidin-4-yl]-L-leucinamide,
- N~1~-(Cyanomethyl)-N~2~{2-[methyl(pyridin-3-ylmethyl)amino]pyrimidin-4-yl}-L-leucinamide,
- N~2~{2-[Benzyl(methyl)amino]pyrimidin-4-yl}-N~1~-(cyanomethyl)-L-leucinamide,
- N~2~{2-[4-(4-Chlorophenyl)piperazin-1-yl]pyrimidin-4-yl}-N~1~-(cyanomethyl)-L-leucinamide,
- N~2~{2-[4-(5-Chloropyridin-2-yl)piperazin-1-yl]pyrimidin-4-yl}-N~1~-(cyanomethyl)-L-leucinamide,

N~1~-(Cyanomethyl)-N~2~-[2-[methyl(thien-3-ylmethyl)amino]pyrimidin-4-yl]-L-leucinamide,

N~1~-(Cyanomethyl)-N~2~-(2-thiomorpholin-4-ylpyrimidin-4-yl)-L-leucinamide,  
N~1~-(Cyanomethyl)-N~2~-[2-(4-phenyl)piperazin-1-yl]pyrimidin-4-yl]-L-leucinamide,  
N~1~-(Cyanomethyl)-N~2~-[2-[2-(hydroxymethyl)piperidin-1-yl]pyrimidin-4-yl]-L-leucinamide,

N~1~-(Cyanomethyl)-N~2~-[2-[(2R)-2-(hydroxymethyl)pyrrolidin-1-yl]pyrimidin-4-yl]-L-leucinamide,

N~1~-(Cyanomethyl)-N~2~-[2-(4-hydroxypiperidin-1-yl)pyrimidin-4-yl]-L-leucinamide,  
N~1~-(Cyanomethyl)-N~2~-[2-[4-(2-furoyl)piperazin-1-yl]pyrimidin-4-yl]-L-leucinamide,

N~2~-[2-[3-(Aminocarbonyl)piperidin-1-yl]pyrimidin-4-yl]-N~1~-(cyanomethyl)-L-leucinamide,

N~1~-(Cyanomethyl)-N~2~-[2-[methyl(2-pyridin-2-ylethyl)amino]pyrimidin-4-yl]-L-leucinamide,

N~2~-[2-(4-Benzylpiperidin-1-yl)pyrimidin-4-yl]-N~1~-(cyanomethyl)-L-leucinamide,  
N~1~-(Cyanomethyl)-N~2~-[2-(4-pyridin-2-ylpiperazin-1-yl)pyrimidin-4-yl]-L-leucinamide,

N~1~-(Cyanomethyl)-N~2~-[2-(4-phenylpiperidin-1-yl)pyrimidin-4-yl]-L-leucinamide,  
N~1~-(Cyanomethyl)-N~2~-[2-[4-(2-hydroxyethyl)piperidin-1-yl]pyrimidin-4-yl]-L-leucinamide,

N~2~-[2-[4-(3-Chlorophenyl)piperazin-1-yl]pyrimidin-4-yl]-N~1~-(cyanomethyl)-L-leucinamide,

N~1~-(Cyanomethyl)-N~2~-[2-(4-phenoxy piperidin-1-yl)pyrimidin-4-yl]-L-leucinamide,  
N~1~-(Cyanomethyl)-N~2~-[2-(3-phenylpyrrolidin-1-yl)pyrimidin-4-yl]-L-leucinamide,  
N~1~-(Cyanomethyl)-N~2~-(2-{methyl[(3-methylisoxazol-5-yl)methyl]amino}pyrimidin-4-yl)-L-leucinamide,

and pharmaceutically acceptable salts thereof.

8. (Canceled)

9. (Previously Presented) A pharmaceutical composition which comprises a compound as defined in claim 1 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable diluent or carrier.

10. (Cancelled)

11. (Withdrawn) A method for treating pain, such as neuropathic pain, in a mammal, such as man, in need of such treatment, which comprises administering to said mammal an effective amount of a compound as defined in claim 1, or a pharmaceutically acceptable salt thereof.

12. (Previously Presented) A pharmaceutical composition which comprises a compound according to claim 7 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable diluent or carrier.

13. (Cancelled)

14. (Withdrawn) A method for treating pain, such as neuropathic pain, in a mammal, such as man, in need of such treatment, which comprises administering to said mammal an effective amount of a compound according to claim 7, or a pharmaceutically acceptable salt thereof.

15. (Withdrawn) A compound according to claim 2 in which  $R^3$  is hydrogen.

16. (Withdrawn) A compound according to claim 2 in which  $R^4$  is hydrogen.

17. (Withdrawn) A compound according to claim 2 in which  $R^5$  is hydrogen or phenyl optionally substituted by  $C_{1-6}$  alkyl or  $C_{1-6}$  alkoxy.

18. (Previously Presented) A compound according to claim 3 in which  $R^3$  is hydrogen.

19. (Previously Presented) A compound according to claim 3 in which  $R^4$  is hydrogen.

20. (Previously Presented) A compound according to claim 3 in which  $R^5$  is hydrogen or phenyl optionally substituted by  $C_{1-6}$  alkyl or  $C_{1-6}$  alkoxy.